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(FILE 'HOME' ENTERED AT 14:15:26 ON 03 MAR 2004)

FILE 'MEDLINE, CAPLUS, BIOSIS, SCISEARCH' ENTERED AT 14:15:46 ON 03 MAR 2004

- 15426 S MEASLES (W) VIRUS
- L2877148 S HETEROLOGOUS (3A) (PROTEIN OR POLYPEPTIDE) OR MARKER
  - 323 S L1 AND L2
- 7292131 S BIOLOGICAL(W) FLUID OR BLOOD OR SERUM OR SALIVA OR SPUTUM L4
- L5 133 S L3 AND L4
- L6 88 DUP REM L5 (45 DUPLICATES REMOVED)
- 7280120 S (SECRET? OR DETECT?) (8A) BIOLOGICAL (W) FLUID OR BLOOD OR SERUM Ь7
- 133 S L3 AND L7 L8

## => d au ti so 50-88 16

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- Scandinavian journal of immunology, (1995 Dec) 42 (6) 581-90. SO Journal code: 0323767. ISSN: 0300-9475.
- MEDLINE on STN ANSWER 49 OF 88 T<sub>1</sub>6
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- Brown D W; Ramsay M E; Richards A F; Miller E ΑU
- Salivary diagnosis of measles: a study of notified cases in the United TIKingdom, 1991-3.
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- OBJECTIVES--To validate a method for salivary diagnosis of measles and to ABassess the diagnostic accuracy of notified cases of measles. DESIGN--Blood and saliva samples were collected within 90 days of onset of symptoms from patients clinically diagnosed as having measles and tested for specific IgM by antibody capture radioimmunoassay. SETTING--17 districts in England and one in southern Ireland during August 1991 to February 1993. SUBJECTS--236 children and adults with measles notified by a general practitioner. RESULTS--Specific IgM was detected in serum in only 85 (36%) of the 236 cases. In cases associated with

outbreaks and tested within six weeks of onset, 53/57 (93%) of samples were IgM positive, thereby confirming the sensitivity of serum IgM detection as a marker of recent infection. The serological confirmation rate was lower in cases with a documented history of vaccination (13/87; 15%) than in those without (70/149; 47%) and varied with age, being lowest in patients under a year, of whom only 4/36 (11%) were confirmed. Measles specific IgM was detected in 71/77 (92%) of adequate saliva samples collected from patients with serum positive for IgM. In cases where measles was not confirmed, 6/101 had rubella specific IgM and 5/132 had human parvovirus B19 specific IgM detected in serum. CONCLUSIONS -- The existing national surveillance system for measles, which relies on clinically diagnosed cases, lacks the precision required for effective disease control. Saliva is a valid alternative to serum for IgM detection, and salivary diagnosis could play a major role in achieving measles elimination. Rubella and parvovirus B19 seem to be responsible for a minority of incorrectly diagnosed cases of measles in the United Kingdom and other infectious causes of measles-like illness need to be sought.

L6 ANSWER 54 OF 88 MEDLINE on STN

DUPLICATE 15

AU Rutter G; Hohenberg H

TI Label-fracture of plasma membranes isolated from measles virus-infected cells.

journal of histochemistry and cytochemistry: official journal of the Histochemistry Society, (1993 Jul) 41 (7) 1085-91.

Journal code: 9815334. ISSN: 0022-1554.

We present a method that permits correlation of the intramembrane AΒ architecture of plasma membrane fracture faces with the distribution of specific molecules at the corresponding cytoplasmic or exoplasmic membrane surfaces. HeLa cells infected with measles virus were used as a model system. Large fragments of the dorsal membrane were isolated after the virus glycoproteins were tagged at the outer cell surface with immune serum and protein A-gold markers. In a second step, different virus polypeptides at the inner cell surface were also identified by a smaller gold label. Thereafter, the isolated plasma membranes were frozen and freeze-fractured. The complementary fracture faces were shadowed with heavy metals and carbon and examined in the transmission electron microscope without cleaning of remaining biological material. Thus, the micromorphology of the replicated fracture faces and the topochemistry of virus components localized at the corresponding leaflets of the plasmalemma could be seen on the same image at high resolution. Of note is that the freeze-fracture morphology of the protoplasmic face is related to the molecular composition of the cytoplasmic surface, as revealed by antibody tagging.

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